Retrograde Ret signaling controls sensory pioneer axon outgrowth

Adam Tuttle¹, Catherine M Drerup¹†, Molly Marra¹,2‡, Hillary McGraw¹§, Alex V Nechiporuk¹*

¹Department of Cell, Developmental, and Cancer Biology, Oregon Health & Science University, Portland, United States; ²Neuroscience Graduate Program, Oregon Health & Science University, Portland, United States

Abstract The trafficking mechanisms and transcriptional targets downstream of long-range neurotrophic factor ligand/receptor signaling that promote axon growth are incompletely understood. Zebrafish carrying a null mutation in a neurotrophic factor receptor, Ret, displayed defects in peripheral sensory axon growth cone morphology and dynamics. Ret receptor was highly enriched in sensory pioneer neurons and Ret51 isoform was required for pioneer axon outgrowth. Loss-of-function of a cargo adaptor, Jip3, partially phenocopied Ret axonal defects, led to accumulation of activated Ret in pioneer growth cones, and reduced retrograde Ret51 transport. Jip3 and Ret51 were also retrogradely co-transported, ultimately suggesting Jip3 is a retrograde adapter of active Ret51. Finally, loss of Ret reduced transcription and growth cone localization of Myosin-X, an initiator of filopodial formation. These results show a specific role for Ret51 in pioneer axon growth, and suggest a critical role for long-range retrograde Ret signaling in regulating growth cone dynamics through downstream transcriptional changes.

Introduction

One major regulator of axon outgrowth and pathfinding is signaling through neurotrophic factors. Of these, the best studied neurotrophic factors are Nerve Growth Factor (NGF) and Brain-derived Growth Factor (BDNF) that signal through the receptors TrkA and TrkB, respectively. NGF and BDNF can have effects locally on axon terminals or signal retrogradely from the axon terminal to the cell body and induce changes in gene expression. NGF/BDNF long-range retrograde signaling is required for survival of sympathetic neurons, where retrograde transport of active endosomal signaling complexes promotes transcription of pro-survival/anti-apoptotic factors (Riccio et al., 1997; Watson et al., 1999; Ye et al., 2003). In addition to survival, retrograde transport of NGF/TrkA promotes sensory axon outgrowth by inducing transcriptional changes in Serum Response Factor (Wickramasinghe et al., 2008), whereas retrograde axonal transport of BDNF/TrkB is required for BDNF-mediated dendritic growth in cortical neurons (Zhou et al., 2012). In general, however, the transcriptional targets downstream of retrograde neurotrophic factor signaling that promote axon growth are still largely unknown.

The neurotrophic factor receptor ‘REarranged during Transfection’ (Ret) and its major ligand, Glial cell line-Derived Neurotrophic Factor (GDNF), have established roles in axon growth, enteric nervous system and kidney development, and cancer progression. Ret receptor has two primary expressed isoforms, Ret9 and Ret51, each with a unique amino acid sequence of the intracellular C-terminus. The two isoforms have different in vivo signaling properties and roles during development. Initial reports indicated Ret9 is required for proper enteric nervous system and kidney development, while Ret51 is dispensable for these processes in murine and zebrafish models (de Graaff et al., 2001; Heanue and Pachnis, 2008; Wong et al., 2005). However, later reports demonstrated...
an important role for Ret51 in murine kidney and enteric nervous system development, suggesting that the two isoforms are partially redundant in these processes (Jain et al., 2006; Jain et al., 2010). These isoforms are also differentially trafficked and susceptible to degradation in both neurons and non-neuronal cells (Richardson et al., 2012; Tsui and Pierchala, 2010). In sympathetic neuron culture, RET9 degrades more slowly than RET51, which is thought to allow retrograde trafficking of RET9 and ultimately promote neuronal survival (Tsui and Pierchala, 2010). In human epithelial cell lines, RET51 is more frequently localized at the plasma membrane, recycled to the plasma membrane after endocytosis, and more rapidly internalized into endosomes after GDNF treatment than RET9 (Crupi et al., 2015; Richardson et al., 2012). Thus, Ret9 and Ret51 have notably different properties and responses to ligand binding even within the same cell.

Ret/GDNF signaling promotes sympathetic (Enomoto et al., 2001), sensory (Honma et al., 2010), and motor (Bonanomi et al., 2012) axon growth and pathfinding. Long-range retrograde RET/GDNF trafficking from distal axons is required for survival of sensory (DRG) neurons and mature sympathetic neurons in rat primary cell cultures (Coupliez and Ibáñez, 2004; Tsui and Pierchala, 2010). In sympathetic neurons, retrograde transport of GDNF ligand from axon terminals to cell bodies has been observed (Coupliez and Ibáñez, 2004; Tomac et al., 1995) but, thus far, retrograde transport of GDNF/Ret in sensory neurons has not. Ret/GDNF signaling can also promote sympathetic axon growth in vitro; however, retrograde signaling/transport is not required for this role over short periods of time (Bodmer et al., 2011; Tsui and Pierchala, 2010). Long-range GDNF/Ret signaling is required for motor neuron extension in developing mouse embryos where loss of GDNF ligand expression in target muscles leads to reduced neuronal expression of Pea3 transcription factor and disrupts neuronal cell body positioning and dendrite patterning (Haase et al., 2002; Vrieseling and Arber, 2006). However, the mechanisms mediating Ret/GDNF retrograde transport and the full extent of the Ret/GDNF downstream transcriptional response mediating axon growth are unknown and in vivo studies of Ret retrograde transport and signaling are sorely lacking (Ito and Enomoto, 2016).

While Ret/GDNF is required for axon extension during development, whether it has a specific role in pioneer axon outgrowth is unknown. Pioneer neurons are the first to extend axons to a particular region or target, acting as a guide and scaffold for ‘follower’ axons. Pioneer neurons are important in the developing CNS and PNS for the initial navigation to appropriate targets, proper follower axon pathfinding, and promoting follower axon outgrowth. Despite their unique role in neurodevelopment, the molecular program regulates their behavior is not well known.

In vivo study of long-range neurotrophic factor signaling, receptor transport, and pioneer axon growth is technically challenging. Zebrafish embryos offer a number of advantages for these studies, including optical accessibility, rapid embryonic development, and amenability to transgenesis and genetic manipulation. In this study, we use the long sensory axons of the zebrafish posterior lateral line ganglion (pLLG) neurons because their planar character, superficial localization, and relatively rapid pioneer axon outgrowth make them uniquely suited for live imaging. The pLL senses water movement through mechanical stimulation of sensory hair cells that transduce this input through the pLL axons innervating them. During extension, 3 to 6 pLL pioneer axons are guided by the pLL primordium (pLLP), a group of cells that migrate along the trunk from 22 to 48 hr post-fertilization (hpf). Pioneer growth cones typically have longer and more elaborate filopodial protrusions compared to followers (Bak and Fraser, 2003; Kim et al., 1991) and this is the case in pLLG pioneer growth cones, which associate extensively with the migrating pLLP (Sato et al., 2010). The pLL primordium expresses GDNF and this ligand production is required for proper pLLG axon outgrowth, but not pLLG neuron specification or survival (Schuster et al., 2010). This allowed us to explore the specific mechanisms of long-range Ret-mediated axon outgrowth in sensory pioneer neurons.

We find that Ret is highly expressed in pioneer neurons and ret51, but not ret9, isoform is required for pioneer axon outgrowth. In the absence of Ret, pioneer axons display reduced growth cone volume and fewer filopodia. We also show that JNK-interacting protein 3 (Jip3) is required for retrograde transport of activated Ret51 from pioneer growth cones. In jip3 mutants, retrograde transport of Ret51 is reduced, leading to accumulation of activated Ret in pioneer growth cones. Additionally, we observe live retrograde, but not anterograde, co-transport of tagged Ret51 and Jip3 in pioneer sensory axons. Finally, we find myosin-X (myo10), a regulator of filopodial formation, is downregulated in ret mutant sensory neurons. Myo10 protein is reduced in ret mutant pioneer growth cones, and loss of Myo10 produces axon extension defects similar to those in ret mutants.
This work describes a novel requirement for Ret51 in axon growth and pioneer neuron development, identifies a novel regulator of phosphorylated Ret51 retrograde transport, and suggests long-range retrograde Ret signaling regulates downstream transcriptional targets which are required for pioneer sensory axon growth cone dynamics.

Results

Ret expression is elevated in pioneer neurons during axon extension

Previous studies showed that ret is expressed in the pLLG neurons and the Ret ligand GDNF, which is expressed by the pLLP, is required for pLLG axon extension but not pLLG neuronal survival (Schuster et al., 2010). Because Ret receptor isoforms Ret9 and Ret51 have distinct properties, we asked which isoform is required for sensory axon extension. We assayed expression with a ret9-specific probe and a probe targeting a shared region of the ret9 and ret51 transcript (denoted ret9+51) in the pLLG during pioneer axon outgrowth (30 hr post-fertilization – hpf) by fluorescence in situ hybridization. Both ret9 and ret9+51 were expressed in the pLLG, with higher levels of both probes detected in the medial portion of the ganglion (Figure 1A,B, see Figure 1—figure supplement 1, panel A for probe target design).

Based on elevated levels of ret mRNA expression in the medial region of the ganglion, which was previously attributed to the location of pioneer neuron cell bodies (Pujol-Martí et al., 2010), we tested whether Ret protein was enriched in extending pioneer neurons. To label cell bodies of the extending pioneer pLLG axons, tails of 40 hpf TgBAC(neurod:EGFP) transgenic zebrafish embryos (hereafter referred to as neurod:EGFP) (Obholzer et al., 2008) were clipped just proximal to the pioneer neuron terminals with rhodamine dextran-soaked scissors (Figure 1C). After 3 hr, embryos were imaged live to note the location of rhodamine-labeled cell bodies in pLLG, then fixed and immunostained with α-Ret primary antibody (for α-Ret antibody validation, see Figure 1—figure supplement 1, panel C). We found that the majority of rhodamine dextran labeled pioneer axons also displayed high levels of Ret protein (Figure 1D’,E’). The dorsal localization of the cell bodies positive for the ret9+51 probe in the same location of the ganglion strongly suggest that Ret receptor is highly expressed in pioneer neurons.

The Ret51 isoform is required for lateral line axon extension

Because Ret is elevated in pioneer neurons during extension, we asked whether Ret activity is required for pioneer axon outgrowth, and, if so, which Ret isoform is necessary for this process. Using the neurod:EGFP transgene to visualize the pLLG axons, we observed that mutants homozygous for the null ret<sup>hu2846</sup> allele (Knight et al., 2011) (which leads to loss of Ret9 and Ret51 isoforms) had disrupted pLLG axon extension, ultimately producing prematurely truncated axons (Figure 1F, G). We observed 97.4% of homozygous ret<sup>hu2846</sup> mutants had premature truncation in one or both pLLG axons by three dpf (n = 89), although we also observed variability in truncation severity between embryos derived from individual adult mating pairs. However, within a group of embryos derived from a single mating pair, ret<sup>hu2846</sup> phenotypes were consistent, as evidenced by the small standard error of the mean of axon truncation in ret<sup>hu2846</sup> mutants in subsequent rescue experiments (Figure 1H,I). Because of pair to pair variability, in our subsequent experiments we compared ret<sup>hu2846</sup> mutant embryos to sibling controls from the same single mating pair (which leads to loss of Ret9 and Ret51 isoforms) (Knight et al., 2011) (which leads to loss of Ret9 and Ret51 isoforms) (Crupi et al., 2015). Injected animals were assayed at three dpf for axon length, measuring the shortest pLLG nerve in the embryo as a conservative measure of axon extension rescue (Figure 1H,I). Comparing un.injected ret<sup>hu2846</sup> mutant embryos to RET isoform mRNA-injected ret<sup>hu2846</sup> mutant siblings we found that RET51, but not RET9, significantly ameliorated the pLLG axon truncation phenotype (ret<sup>hu2846</sup> mutant uninjected mean body segment location of axon termination: 17.5 ± 1.7 vs. ret<sup>hu2846</sup> mutant RET9 mRNA injected: 18.1 ± 1.6, p=0.001 by one-way ANOVA with post-hoc Tukey test, see for further details; ret<sup>hu2846</sup> mutant uninjected: 13.8 ± 0.7 vs. ret<sup>hu2846</sup> mutant RET51 injected: 18.8 ± 0.9, p<0.001). In some cases, RET51 injection produced full-length, bilateral rescue of axon extension (Figure 1I, 9 of n = 87 embryos). This suggests that Ret51 is the critical isoform required for Ret-mediated pLLG axon outgrowth.
Figure 1. ret is highly expressed in pLL pioneer neurons and required for pLL axon outgrowth. (A,B) Lateral view of ret9 and ret9+51 probes, visualized by fluorescence in situ hybridization in neurod:EGFP embryos, reveals both are expressed mostly in dorsal cells during axon extension (30 hpf) in the pLLG (yellow line = pLLG outline, scale bar = 20 µm; probe design schematized in Figure 1—figure supplement 1A). (C) Diagram of pLL nerve sever Figure 1 continued on next page
Ret is necessary for proper pioneer growth cone morphology

Because pLLG axon extension is incomplete in ret\textsuperscript{hu2846} mutants, we examined the cellular bases of this axon growth failure. After initial axon outgrowth and premature termination of axon extension in ret\textsuperscript{hu2846} mutant embryos, we observed no degeneration or retraction of the partially extended pLLG axons from 48 hpf through 120 hpf (Figure 1—figure supplement 2, panel A). To determine if there was a difference in the number of pioneer neurons that extended axons during initial outgrowth, we immunostained ret\textsuperscript{hu2846} mutants and their wild-type siblings with α-SCG10 antibody at 24 (early outgrowth) or 30 hpf (mid-outgrowth) and imaged distal pLLG axons. SCG10 (STMN2) is enriched in extending sensory pioneer growth cones (Drerup et al., 2016), serving as a marker of individual pioneer growth cones. Based on SCG10 expression in the distal 100 μm of axons (visualized by the neurod:EGFP transgene), the number of pioneer growth cones that initially extended the pLLG axon was not significantly different between ret\textsuperscript{hu2846} mutants and siblings (Figure 1—figure supplement 2, panel C, wild type: 5.73 ± 0.29 vs. ret\textsuperscript{hu2846}: 5.56 ± 0.48, p=0.99 by Mann-Whitney U test), indicating there is no defect in pioneer neuron specification or initial axon outgrowth in ret\textsuperscript{hu2846} mutants. However, at 30 hpf, ret\textsuperscript{hu2846} mutants had significantly fewer growth cones in the distal 100 μm of the growing pioneer axon bundle (Figure 1—figure supplement 2, panel D, wild type: 6.58 ± 0.32 vs. ret\textsuperscript{hu2846}: 4.00 ± 0.44, p<0.001). These observations suggest that pLLG pioneer neurons are specified and begin to extend normally; however, axon extension fails after initial outgrowth.

To explore the effect of Ret loss-of-function on pioneer growth cone morphology and behavior, we assayed axon terminals of extending pioneer axon bundles in ret\textsuperscript{hu2846} mutants and wild-type siblings (Figure 2A,B). First, we quantified axonal volume of the distal 75 μm of neurod:EGFP transgenic embryos at 30 hpf which typically contains 5–6 pioneer growth cones (Figure 2C). In comparison to siblings, ret\textsuperscript{hu2846} mutants had significantly reduced distal axonal volume in extending pioneer axon bundles (Figure 2C; wild type: 1477.6 ± 48.9 μm\textsuperscript{3} vs ret\textsuperscript{hu2846}: 1137.4 ± 136.1, p=0.003). To visualize individual pioneer neuron and growth cone morphology, neurod:EGFP embryos at the one-cell stage were injected with plasmid containing the neurod5kb promoter (Mo and Nicolson, 2011) driving mCherry (neurod5kb:mCherry). Injected plasmid distributes mosaically in zebrafish embryos, allowing identification and live imaging of individual pLLG pioneer growth cones expressing mCherry during extension (30 hpf) (Figure 2A’,B’). Compared to wild-type siblings,
ret<sup>hu2846</sup> mutant pioneer growth cones were much less elaborate, had significantly reduced volume (Figure 2D; wild type: 291.5 ± 24.2 µm<sup>3</sup> vs. ret<sup>hu2846</sup>: 182.6 ± 17.4, p<0.001), and significantly fewer filopodia (defined as protrusions ≥ 1 µm) per growth cone (Figure 2E, wild type: 9.4 ± 0.7 vs. ret<sup>hu2846</sup>: 5.9 ± 0.6, p<0.001). However, the length of these filopodia was not significantly different (Figure 2F, wild type: 4.55 ± 0.22 µm vs. ret<sup>hu2846</sup>: 4.08 ± 0.29, p=0.20) between the two groups. Together, these data show that Ret signaling is required for maintaining proper pioneer growth cone morphology and filopodial numbers but not length.

**Jip3 loss-of-function phenocopies loss of Ret signaling**

Neurotrophic factor signaling often involves retrograde transport of the activated receptor-ligand complex from the axon terminal to the cell body. However, the specific trafficking mechanisms are not clearly understood. In the pLLG, JNK-interacting protein 3 (Jip3) functions as a specific retrograde cargo adaptor for Dynein-mediated axonal transport during development (Drerup and Nechiporuk, 2013). Embryos homozygous for the Jip3 null allele, jip3<sup>nl7</sup>, display pLLG axon extension defects similar to ret<sup>hu2846</sup> mutants (Figure 3B) (Drerup and Nechiporuk, 2013). Compared to siblings, jip3<sup>nl7</sup> mutant pLL nerves were significantly truncated by three dpf (Figure 3—figure...
Figure 3. Loss of Jip3 phenocopies ret mutant pioneer axon defects. (A,B) Lateral view, live image of wild-type sibling (A) and jip3<sup>n17</sup> mutant (B) containing the neurod:EGFP transgene at two dpf. Jip3<sup>n17</sup> mutants display pLLG axon truncation defects (arrowhead; mutant truncation phenotype plotted in Figure 3—figure supplement 1). (C, D) Lateral view, live image of collective pioneer axon terminals of wild-type siblings (C) and jip3<sup>n17</sup> mutants (D). Figure 3 continued on next page.
Figure 3 continued

labeled by neurod:EGFP transgene at 30 hpf. Scale bar = 20 μm (C–D). Individual pioneer axons in the same animal labeled by mosaic expression of neurod5kb:mCherr plasmid: (E–F) Quantification of distal collective axon terminal volume (E) or individual pioneer growth cone volume (F). jip3\(^{\text{nl7}}\) mutants have significantly reduced collective axonal volume but not individual growth cone volume. (G–H) Quantification of number (G) and length (H) of filopodia (>1 μm) per individual pioneer axon. Jip3\(^{\text{nl7}}\) mutants have significantly reduced number of filopodia but not length. (I) Quantification of axon truncation defects of sibling, heterozygous, and homozygous jip3\(^{\text{nl7}}\) mutants in homozygous ret\(^{\text{hu2846}}\) background shows no significant difference in axon truncation phenotype between genotypes. Jip3\(^{\text{nl7}}\) homozygous mutants with +/- or +/-ret\(^{\text{hu2846}}\) background had significantly less severe axon truncations. Error bars represent S.E.M., *=p < 0.05, N.S. = not significant.

DOI: https://doi.org/10.7554/eLife.46092.008

The following source data and figure supplement are available for figure 3:

**Source data 1.** Individual embryo scoring data for scoring of growth cone and filopodia metrics in jip3 and axon truncation in jip3/ret double mutants.

DOI: https://doi.org/10.7554/eLife.46092.010

**Figure supplement 1.** Quantification of jip3\(^{\text{nl7}}\) homozygous mutant axon truncation defects.

DOI: https://doi.org/10.7554/eLife.46092.009

supplement 1; mean body segment location of termination sibling: 33 ± 0 vs. jip3 mutants: 24.0 ± 0.6, p<0.0001 by Mann-Whitney U test). Thus, we investigated whether Jip3 and Ret act in the same pathway during axon outgrowth. We discovered that, similar to ret\(^{\text{hu2846}}\) mutants, jip3\(^{\text{nl7}}\) mutants displayed reduced terminal collective pioneer axon volume at 30 hpf (Figure 3D,E, wild type: 1294.7 ± 51.5 μm\(^3\) vs jip3\(^{\text{nl7}}\): 1134.7 ± 57.2, p=0.024) and fewer filopodia per pioneer growth cone (Figure 3G, wild type: 10.3 ± 0.9 vs. jip3\(^{\text{nl7}}\): 7.7 ± 0.4, p=0.017) with no significant change in filopodial length (Figure 3H, wild type: 3.88 ± 0.38 μm vs. jip3\(^{\text{nl7}}\) mutant: 4.24 ± 0.26, p=0.33).

Additionally, double homozygous ret\(^{\text{hu2846}}\) mutant and jip3\(^{\text{nl7}}\) mutant had no significant difference in axon truncation defects compared to single homozygous ret mutants (Figure 3I, body segment location of axon termination in ret\(^{\text{hu2846}}\) mutants, +/-: 14.4 ± 1.7, jip3\(^{\text{nl7}}\)/+: 13.9 ± 3.0, jip3\(^{\text{nl7}}\)/jip3\(^{\text{nl7}}\): 15.7 ± 3.9, p=0.92 by one-way ANOVA), indicating Jip3 and Ret operate in the same genetic pathway. Notably, wild-type ret siblings homozygous for jip3\(^{\text{nl7}}\) had a less severe axon truncation defect compared to the jip3/ret homozygous double mutants (jip3\(^{\text{nl7}}\)/jip3\(^{\text{nl7}}\), +/- or ret\(^{\text{hu2846}}\)/+: nl7 ± 1.1; p=0.047 by post hoc Tukey test). Based on these findings, we investigated whether loss of Jip3 affects localization of Ret protein in the extending axons. jip3\(^{\text{nl7}}\) mutants displayed significant accumulation of both total and phospho-Y905 (activated) Ret receptor (referred to as tRet and pRet, respectively; for α-Ret antibody validation, see Figure 1—figure supplement 1, panels B-E) at the pioneer growth cone during extension compared to wild-type siblings (Figure 4A–F, for tRet, wild type fluorescence intensity: 1.97 ± 1.15 vs. jip3\(^{\text{nl7}}\): 45.09 ± 12.44, p<0.001; for pRet, wild type: 7.59 ± 1.60 vs. jip3\(^{\text{nl7}}\): 20.54 ± 3.90, p=0.033). These data suggest that a pool of activated Ret receptor fails to be trafficked from the extending growth cone in jip3\(^{\text{nl7}}\) mutants.

We next asked whether retrograde movement of Jip3 is necessary for the clearance of Ret from the growth cone and axon extension. To test this, we injected EGFP-tagged mRNAs encoding full-length Jip3 (control), Jip3 lacking the p150 binding domain that is unable to interact with the Dynnein motor complex (Cavalli et al., 2005) (jip3\(Δ\)p150) or Jip3 lacking the JNK interacting domain (jip3\(Δ\)JNK) (Morfini et al., 2009) into one-cell stage embryos from a heterozygous +/-/+ret\(^{\text{hu2846}}\) mutant incross (Figure 4G). Embryos were fixed at 3 days and immunostained for neurofilaments (3A10) to visualize pLLG axons and the proportion of embryos with axon truncation were scored. Expression of a full-length jip3 completely rescued axon truncation at 3 dpf (Figure 4G, uninjected truncation rate: 17.4%, n = 23 vs. jip3: 0%, n = 29). Within embryos from a single mating pair, jip3\(Δ\)JNK also rescued truncation while jip3\(Δ\)p150 failed to rescue (uninjected: 15.8%, n = 38, jip3\(Δ\)JNK: 0%, n = 15, jip3\(Δ\)p150: 22.2%, n = 36). Additionally, expression of full-length jip3 mRNA rescued accumulation of pRet in pioneer growth cones (Figure 4I) but jip3\(Δ\)p150 did not (Figure 4J). Our previous work showed that loss of Jip3 does not affect JNK signaling in extending pioneer pLLG axon growth cones. Similarly, we found no significant difference in pJNK levels in growth cones of ret\(^{\text{hu2846}}\) mutants and wild-type siblings at 30 hpf (Figure 4—figure supplement 1). Altogether, these data indicate that the Jip3 association with the retrograde motor is required for Jip3-
Figure 4. Jip3 is required for clearance of activated Ret from pioneer axon growth cones. (A–D) Lateral view, immunostained pioneer axon growth cones labeled by neurod:EGFP transgene at 30 hpf. Immunostaining for total Ret receptor (A',B', WT n = 8, jip3<sup>nl7</sup> n = 5) and activated pRet905 receptor (C',D', WT n = 4, jip3<sup>nl7</sup> n = 4) shows no signal in wild-type sibling pioneer growth cones (A',C') but a notable accumulation of total and pRet905.
dependent axon extension and clearance of pRet from pLLG growth cones, suggesting disrupted Ret retrograde transport may underlie the axon truncation observed in jip3<sup>−/−</sup> mutants.

**Jip3 mediates retrograde transport of activated Ret51 receptor**

Based on the role of Jip3 as a retrograde transport adapter and the similarity of ret<sup>hu2846</sup> and jip3<sup>−/−</sup> mutant axonal phenotypes, we asked if Jip3 mediates retrograde transport of Ret. To address this question, we first performed nerve sever experiments (Drerup and Nenichporuk, 2013) as an indirect readout of Ret axonal transport (Figure 5A–F). Post nerve sever, anterogradely transported cargoes accumulate in the proximal stump, whereas retrogradely transported cargoes accumulate in the distal stump (Drerup and Nenichporuk, 2013). We hypothesized that if Jip3 specifically mediates retrograde axonal transport of Ret from the growth cone, Ret protein should accumulate at the distal site of nerve sever. Extended pLLG pioneer axons (labeled by neurod:EGFP) of siblings and jip3<sup>−/−</sup> mutants were cut between neuromast 2 and 3 at five dpf, allowed to recover for 3 hr, and immunostained for tRet and pRet. Total Ret was detected at the proximal and distal cut sites, indicating that Ret is bidirectionally trafficked in axons. However, the amount of total Ret was not different between wild-type and jip3<sup>−/−</sup> mutant embryos (Figure 5A–D), proximal fluorescence intensity, wild type: 32.0 ± 2.5 vs. jip3<sup>−/−</sup>: 28.4 ± 3.5, p=0.39; distal FI, wild type: 26.2 ± 2.3 vs. jip3<sup>−/−</sup>: 28.4 ± 2.8, p=0.54), arguing that Jip3 is not required for this process. In wild-type siblings and jip3<sup>−/−</sup> mutant embryos, we observed no accumulation of pRet at the proximal stump (Figure 5C,F, wild type: 9.0 ± 2.0 vs. jip3<sup>−/−</sup>: 9.8 ± 3.1, p=0.82); however, pRet accumulated significantly at the distal stump of wild-type siblings (Figure 5C) compared to jip3<sup>−/−</sup> mutants, which had much less pRet accumulation at the distal cut site (Figure 5D',F, wild type: 22.4 ± 2.9 vs. jip3<sup>−/−</sup>: 11.4 ± 1.8, p=0.004). Altogether, these data imply that Jip3 is specifically required for retrograde transport of activated Ret.

Next, we tested directly if Jip3 is required for axonal transport of Ret receptor by observing trafficking of Ret fusion protein in extending pLLG axons. To achieve this, we injected a plasmid driving mCherry-tagged RET51 under the neurod5kb promoter (neurod5kb:RET51-mCherry) at the one-cell stage and imaged RET51 transport in extending pLLG axons at 30 hpf (Drerup and Nenichporuk, 2016). Injected plasmid distributes mosaically in the developing embryo, thus, we can select embryos expressing fusion constructs in 1–3 pLLG neurons to assay Ret trafficking in individual axons during extension. We observed anterograde and retrograde Ret trafficking in embryos injected with neurod5kb:RET51-mCherry or ret51-EGFP (Figure 5—figure supplement 1). To test whether Jip3 regulates Ret51 retrograde axonal transport in pioneer neurons, we assayed RET51-mCherry transport in jip3<sup>−/−</sup> embryos and wild-type siblings. For our experiments, we used expression of the neurod:EGFP transgene to identify the most distal pLLG pioneer neurons also expressing RET51-mCherry (Figure 5G,H). Based on kymograph analysis, we found significantly fewer retrogradely trafficked RET51-mCherry particles in jip3<sup>−/−</sup> mutant pLLG pioneer neurons compared to wild-type
Figure 5. Jip3 mediates retrograde axonal transport of activated Ret51. Lateral view, immunostaining of severed pLL nerves proximal (A–D) and distal (A’–D’) to injury site, 3 hr post-injury; dotted line = nerve outline. (A,B) tRet immunostaining is not notably different between wild-type siblings (A,A’) and jip3^nl7 mutants (B,B’) on either side of injury site. (C,D) p905Ret staining is mostly absent in wild-type siblings proximal to injury site (C) and jip3^nl7 mutants on both sides of the injury site (D,D’). Scale bar = 10 μm. Stronger p905Ret staining signal is present in wild-type siblings distal to the injury.
siblings, suggesting a disruption in retrograde transport of Ret51 (Figure 5G',H',L). To examine whether transport of other neurotrophic factor receptors expressed in the pLLG, such as TrkB (Gasanov et al., 2015), was disrupted, we analyzed trafficking of TrkB-mCherry fusion from embryos injected with neurod5kb:trkB-mCherry plasmid (Figure 5I,J). Notably, we saw no significant change in trafficking of TrkB-mCherry particles (Figure 5K,L). These data, taken together with pRet accumulation in the distal stump of severed jip3<sup>nu7</sup> mutant pLLG axons, strongly argue that Jip3 is required for transport of activated Ret51, but not TrkB, in pLLG axons.

Finally, if Jip3 acts as a specific cargo adaptor for retrograde pRet51 transport, we expect that Jip3 and Ret51 would be co-transported in extending pLLG axons. To address this, we co-injected Tg(BAC)Ret51-EGFP and neurod5kb:jip3-mCherry (Drerup and Nechohiruk, 2013) plasmid at one-cell stage and imaged pLLG pioneer axons expressing both fusions at 30 hpf. Co-localization of Jip3-mCherry and Ret51-EGFP was observed in a subset of retrogradely trafficked particles but no cases of co-localization were observed for anterogradely transported particles (Figure 6A,B, Figure 6—video 1). Taken together, these data suggest Jip3 preferentially mediates retrograde transport of activated pRet51.

Because Jip3 mediates retrograde transport of pRet and jip3<sup>nu7</sup> mutants display axonal phenotypes that are similar to the ret<sup>hu2846</sup> mutant, we hypothesized that Jip3 mediates long-range Ret signaling which is required for pLLG axon outgrowth. However, jip3<sup>nu7</sup> mutants typically display less severe pLL axon truncations than ret<sup>hu2846</sup> mutants (Figures 1I and 3I, S3). We reasoned that this difference may be due to both local and long-range requirements for Ret signaling in pLLG axons. To distinguish between the local and long-range signaling, we looked for ways to acutely inactivate Ret receptor in jip3<sup>nu7</sup> mutants. To achieve this, we used a multi-kinase inhibitor, sorafenib, which blocks signaling of multiple receptor-tyrosine kinases including Ret, where it potently inhibits Ret activation (Plaza-Menacho et al., 2007). To confirm that sorafenib can inhibit Ret in pLLG axons, we treated ret<sup>hu2846</sup> mutant and wild-type sibling embryos with 5 μM sorafenib from 24 to 72 hpf (Figure 6—figure supplement 1). Sorafenib treatment of wild-type siblings induced pLLG axon defects similar to those found in untreated ret mutants. Furthermore, we observed no significant exacerbation of the axon truncation in ret mutants treated with sorafenib compared to either untreated ret mutants or sorafenib-treated siblings. This suggests that, in this context, sorafenib functions as a specific inhibitor of Ret signaling. We then treated jip3 mutant and sibling larvae with sorafenib (Figure 6C). Drug treatment induced significantly more severe axon truncations in siblings compared to untreated jip3 mutants (mean body segment location of axon termination in sorafenib-treated siblings: 16.1 ± 1.5 vs. untreated jip3<sup>nu7</sup> mutants: 22.5 ± 1.1, p<0.001 by one-way ANOVA with post-hoc Tukey test). In addition, sorafenib-treated jip3 mutants displayed similar truncation severity to sorafenib-treated siblings (sorafenib-treated jip3<sup>nu7</sup> mutants: 15.4 ± 1.0, p=0.96). These results are consistent with the observed difference in axon truncation severity between ret and jip3 mutants and suggest local Ret signaling is required for long-range Ret signaling.
signaling in the growth cone promotes some aspects of axon outgrowth even in the absence of Jip3-mediated long-range retrograde transport.

**Ret signaling regulates transcription and pioneer growth cone levels of Myosin-X**

Because retrograde transport is required for some aspects of Ret-mediated pLL sensory axon growth, we asked if there were specific transcriptional targets downstream of long-range Ret signaling that are required for pioneer axon outgrowth. Based on the observed defects in filopodial formation and growth cone advancement, we looked for factors that are specifically expressed in pLLG...
neurons during axon extension and regulate filopodial dynamics. We focused on Myosin-X (Myo10), an unconventional Myosin that promotes initiation of filopodia and links integrins and the actin cytoskeleton to promote cell migration. Myo10 is upregulated during neurite growth following peripheral nerve injury (Plantman et al., 2013) and loss of Myo10 disrupts cortical axon growth (Zhu et al., 2007). Although humans and mice have one Myo10 gene, zebrafish have three with specific expression patterns. Of these, Myo10l1 has the most restricted embryonic expression and is the only one expressed in the pLLG (Sittaramane and Chandrasekhar, 2008). We hypothesized that loss of Ret signaling would reduce myo10l1 transcription, leading to decreased Myo10 in pioneer growth cones. In turn, this would disrupt filopodia formation and subsequent axon growth in ret mutants. To test this, we analyzed myo10l1 levels in rethu2846 mutants and siblings at 30 hpf using fluorescence in situ hybridization (Figure 7A,B). rethu2846 mutant ganglia had significantly reduced mean fluorescence intensity compared to siblings (Figure 7B,C). To verify if this reduced Myo10 protein in pioneer growth cones, we immunostained 30 hpf rethu2846 mutants and sibling neurod:EGFP embryos with Myo10 antibody (Figure 7D,E). This antibody recognized Myo10l1-eGFP fusion protein expressed in zebrafish sensory neurons using BAC transgenesis (Figure 7—figure supplement 1, panel A). Consistent with the observed changes in myo10l1 transcription, we found significantly reduced Myo10 immunostaining fluorescence in distal pioneer axons (Figure 7E,F). Similarly, the levels of Myo10l1 were reduced in axon growth cones of jip3 mutants (Figure 7G). If Myo10l1 is required for proper pioneer axon outgrowth, we expect loss of Myo10l1 to disrupt pLLG axon growth, similar to loss of Ret signaling. To test this, we performed G0 CRISPR-Cas9-mediated genetic knockdown of myo10l1 using the established method of injecting a multiplexed cocktail of three sgRNAs (Shah et al., 2015; Wu et al., 2018) that efficiently target three different regions of myo10l1 in developing neurod: EGFP embryos (Figure 7—figure supplement 1, panels B,C). Overall, myo10l1 crispant embryos displayed significant pLL nerve truncation compared to uninjected siblings (Figure 7H, crispant mean pLL truncation: 27.4 ± 1.0 vs. uninjected siblings: 32.9 ± 0.1, p<0.001 by one-way ANOVA with post-hoc Tukey test, see Materials and methods for more details; n = 112 and 136, respectively), although pLL truncation was partially penetrant (n = 34 of 112 embryos). This suggests Myo10l1 is required for proper pioneer axon outgrowth and loss/reduction of Myo10l1 may contribute to rethu2846 mutant axon truncation defects. Reduction of Myo10l1 in rethu2846 mutants did not exacerbate the axon truncation phenotype (17.0 ± 1.2 vs. myo10l1 crispant rethu2846 mutants: 17.1 ± 1.7, p=1.00; Figure 7G; n = 36 and 29, respectively), suggesting that Ret and Myo10l1 function in the same pathway.

Discussion

While the ability of GDNF/Ret signaling to promote axon outgrowth during development is known, the mechanisms of long-range retrograde Ret signaling and consequent effects on transcription and pioneer axon behavior are largely unknown. Our work finds that the Ret51 isoform is required for sensory pioneer axon outgrowth. We observed enriched expression of ret9+51 mRNA in a region of the pLLG known to hold the pioneer neuron cell bodies (Pujol-Martí et al., 2010). Through retrograde dye labeling of pioneer neuron cell bodies, we found an enrichment of Ret protein in pLLG pioneer neuron cell bodies, corresponding to the region of ret9+51 mRNA expression. rethu2846 mutants do not have defects in pLLG pioneer neuron specification or survival but do have a failure of continued growth of pioneer axons. This pioneer axon growth was rescued in rethu2846 mutants by exogenous expression of RET51 mRNA but not RET9, indicating that Ret51 is specifically required for pioneer axon outgrowth. These data also demonstrate that Ret is a specific marker of pioneer neurons in the pLLG.

We propose that long-range retrograde Ret signaling mediated by Jip3 is required for pioneer axon outgrowth and growth cone dynamics. While Ret/GDNF signaling could have local effects on growth cone behavior and morphology, we argue that long-range retrograde signaling and a transcriptional response underlie the loss of filopodia and a significant portion of the pLLG axon truncation phenotype described for several reasons. First, the Jip3 and Ret loss-of-function axon truncation and growth cone filopodia phenotypes are very similar. Second, there is no difference in the severity of axon truncation in jip3hu2846/rethu2846 double mutants compared to individual ret homozygous mutants, suggesting they act in the same pathway. Third, in jip3 mutants we observe an accumulation of phospho-Ret in extending growth cones and a reduction in retrograde axonal transport of
Figure 7. Myosin-X expression and growth cone localization are reduced in ret<sup>hu2846</sup> mutants. (A,B) Lateral view, pLL ganglia marked by neurod:EGFP of wild type and ret<sup>hu2846</sup> mutants at 30 hpf. (A’,B’) Fluorescent in situ hybridization shows decreased expression of myosin-10l1 in ret mutants (yellow dotted line, ganglion; sc and white dotted line, spinal cord). (C) Quantification of mean fluorescence intensity of fluorescent in situ hybridization reveals significantly lower expression of myosin-10l1 in ret mutant pLL ganglia. (D,E) pLL pioneer axon growth cones at 30hpf marked by neurod:EGFP expression. (D’,E’) Immunostaining for Myo10 shows a decrease in overall fluorescence and puncta (magenta arrowheads) in ret<sup>hu2846</sup> mutants (Antibody confirmation shown in Figure 7—figure supplement 1, panel A). (F,G) Quantification of mean fluorescence intensity of Myo10 immunostaining in the distal 50 μm of pLL pioneer axons shows a significant decrease in Myo10 localization in ret<sup>hu2846</sup> (F) and jip3<sup>nl7</sup> mutants (G). (H) Whisker plot of sibling and ret mutant larvae (neurod:EGFP) injected with myo10l1 gRNA/Cas9. Myo10l1 CRISPR injection in wild-type ret siblings-induced axon truncation in many cases but did not significantly exacerbate axon truncation in ret mutants (myo10l1 transcript and CRISPR targeting shown in Figure 7—figure supplement 1, panels B,C). p. Error bars represent mean S.E.M., *p < 0.05, ***p < 0.001 (one-way ANOVA with post hoc Tukey test). (I) Proposed model of the contribution of long-range retrograde Ret signaling to axon outgrowth of pioneer sensory neurons.

DOI: https://doi.org/10.7554/eLife.46092.021

The following source data and figure supplement are available for figure 7:

Figure 7 continued on next page
Ret51 and pRet by in vivo imaging and immunostaining following nerve sever, respectively. Fourth, jip3 constructs lacking the p150 Dynein complex binding domain cannot rescue axon truncation or distal pRet accumulation in jip3 mutants. However, jip3 lacking the JNK binding domain fully rescues these defects, and we observe no changes in growth cone pJNK levels in jip3 or ret mutants, indicating Jip3 retrograde transport is required for proper axon outgrowth and retrograde transport of pRet from extending growth cones. Finally, in ret mutants we observe a reduction in myo10l1 expression and Myo10l1 in pioneer growth cones, suggesting a specific downstream transcriptional target of Ret signaling that underlies one of the observed growth cone defects. Consistent with this, knockdown of Myo10l1 disrupted pioneer axon extension similar to rethu2846 mutants. Thus, we provide evidence for a mechanism by which long-range retrograde signaling induces a specific transcriptional change that in turn regulates pioneer growth cone dynamics and outgrowth. This model is consistent with previous observations in cultured sympathetic and sensory neuron indicating that neurotrophic factor-dependent axon growth over long periods of times requires a transcriptional response (Bodmer et al., 2011).

Our data also provide support that, in addition to a long-range Ret-mediated retrograde signaling, local Ret activity is necessary for pLL axon growth. We found that genetic loss of Jip3, which presumably blocks retrograde Ret signaling, produces a less severe axon truncation phenotype than loss of Ret. We also find that pharmacological inhibition of Ret in wild-type axons induces a more significant axon truncation than in untreated jip3 mutants. Finally, we noted differences in the pLLG pioneer axons of jip3 mutants compared to ret mutants. Specifically, unlike ret mutants, individual growth cone volume is not significantly reduced in jip3 mutants. Overall, these observations argue that a combination of both local and long-range Ret signaling are required for axon outgrowth. In fact, we would like to suggest that Jip3-mediated retrograde transport may not be necessary for early stages of the pLLG pioneer axon outgrowth, but is required for continued axon growth as axons extend beyond a certain distance from the soma. This model is consistent with previous observations in cultured sympathetic and dorsal rood ganglia neurons, in which long-range retrograde transport of TrkA receptor is required for long-term (but not short-term) maintenance of axon outgrowth (Bodmer et al., 2011).

**Ret signaling regulates pioneer growth cone morphology and dynamics**

Regulated growth cone dynamics, including formation and disassembly of filopodia and lamellipodia, is required for growth cone advancement and axon outgrowth. Loss of Ret led to fewer filopodia per pioneer growth cone but did not change mean length of filopodia. This suggests a defect in initiation of filopodial formation but not maintenance or continued growth. Consistent with this result, we observed reduction in Myo10l1, an unconventional Myosin that promotes filopodia formation, in ret mutants (Bohil et al., 2006; Zhu et al., 2007). In addition, transient CRISPR-based knockdown of Myo10l1 in ret mutants did not exacerbate the axon truncation phenotype observed in ret mutants alone, suggesting that Myo10l1 acts downstream of Ret. Although, to conclusively determine if myo10l1 acts in a local pathway with Ret in pLLG neurons during pioneer axon extension, germline ret;myo10l1 double mutants carrying the neurod:EGFP transgene should be generated and their offspring analyzed for axon truncation defects. Disruption of dynamic filopodial formation can prevent proper growth cone motility (Lowery and Van Vactor, 2009), leading to failed axon outgrowth. Another non-mutually exclusive possibility is that failure to dynamically form filopodia could disrupt proper interaction with the pLLP (which produces GDNF and BDNF; Gasanov et al., 2015) necessary for guiding and promoting axon growth. In this case, less interaction with the pLLP could reduce reception of trophic/permissive cues produced by the pLLP, further reducing the formation of growth cone filopodia and producing a positive feedback cycle that ultimately ends in failed growth cone advancement.
In addition to the loss of filopodia, we also observe a significant reduction in pioneer growth cone volume. The reduced growth cone volume and more compact morphology of ret mutant growth cones which lack the typical spread out ‘footprint’ of pioneer growth cones could be caused by defects in formation of lamellipodia, which would ultimately prevent proper growth cone migration and steering. While changes in myo10 expression are consistent with the filopodial defects and axon outgrowth failure, local Ret signaling may promote lamellipodia formation or other specific growth cone behaviors.

**Jip3 is an adapter for transport of activated Ret**

While there is evidence that long distance GDNF signaling is required for motor axon pathfinding (Vrieseling and Arber, 2006), to date there has been no in vivo observation of retrograde axonal trafficking of Ret receptor or description of the mechanisms that would underlie this transport in neurons (Ito and Enomoto, 2016). In this study, we provide compelling evidence that Jip3 functions as an adapter for the retrograde transport of activated Ret to promote axon outgrowth. We found that Jip3 loss-of-function partially phenocopied pLLG axon truncation and many of the growth cone defects we observed in ret<sup>hu2846</sup> mutants. We observed no difference in axon truncation defects in jip3<sup>nl7</sup> and ret<sup>hu2846</sup> null double mutants compared to ret<sup>hu2846</sup> mutants, indicating that Ret and Jip3 act in the same pathway to promote axon growth. Our data also show that loss of Jip3 leads to accumulation of activated Ret in pioneer axon growth cones. The Jip3-mediated clearance of pRet requires Jip3 binding to the retrograde motor, consistent with the idea that Jip3 functions as a retrograde adapter in this context. Axon sever experiments showed that activated Ret is retrogradely transported in axons and Jip3 is required for that process. Consistent with this observation, we found a significant reduction in the number of retrogradely trafficked Ret51 particles in jip3<sup>nl7</sup> mutants. Notably, we did not observe complete loss of retrograde transport of Ret51 in jip3<sup>nl7</sup> mutants or observe complete co-localization of Jip3/Ret51 retrograde transport. This is likely due to the inability to distinguish between phosphorylated or non-activated tagged Ret receptor in vivo. These observations are consistent with our assumption that Jip3 is required for retrograde transport of activated Ret51, which presumably constitutes only a subset of Ret51 fusion visualized in our live trafficking experiments. Finally, we observed live retrograde co-transport of Jip3 and Ret51 tagged particles in extending pLLG axons, but no co-transport of anterogradely trafficked particles. Collectively, these results indicate Jip3 mediates retrograde transport of phosphorylated Ret51 in pioneer axons.

It is possible that Jip3 mediates local Ret signaling in pioneer growth cones, since it also can function as a scaffold that regulates local JNK phosphorylation. However, while Jip3 interaction with JNK has been previously implicated in axon extension (Sun et al., 2013), it is not necessary for pLL axon extension (Dreerup and Nchiporuk, 2013) and we observe no change in pJNK levels in pioneer growth cones in ret<sup>hu2846</sup> mutants. Consistent with this, we found that mRNA injection of full-length Jip3 and Jip3 lacking the JNK-binding domain in jip3<sup>nl7</sup> mutants completely rescued axon truncation and accumulation of pRet in pioneer growth cones. Combined with our previous results, this indicates Jip3-mediated retrograde axonal transport of phosphorylated Ret51 is required for axon outgrowth and proper formation of growth cone filopodia.

**A specific role for Ret51 isoform in pioneer axon outgrowth**

Ret receptor isoforms have different roles and requirements during development. Ret9 previously was described as the critical isoform in mouse kidney organogenesis, enteric nervous system development, and embryogenesis in general (de Graaff et al., 2001; Wong et al., 2005). However, later murine studies argue these isoforms have largely redundant developmental function (Jain et al., 2006; Jain et al., 2010). Ret isoform roles are little studied in zebrafish, though initial work indicated Ret9, but not Ret51, is required for proper enteric nervous system development (Heanue and Pachnis, 2008). Neuronal studies examining differences in Ret9 and Ret51 subcellular localization and degradation have focused on their roles in neuronal survival (Tsui and Pierchala, 2010). RET51 is expressed in a specific subset of RET9-expressing mouse olfactory neurons during embryonic development (Kaplinovsky and Cunningham, 2011), but the functional consequence of this is unknown. Our work establishes a novel, specific role for Ret51 in pioneer axon extension unrelated to neuronal differentiation or survival and describes in vivo observation of axonal Ret51 isoform trafficking.
Though Ret9 failed to rescue mutant axon truncation, it may be that Ret9 has a role in local signaling at the dendrites or cell body that is unrelated to axon outgrowth.

These isoform-specific functional differences may be explained by differential trafficking properties and degradation rates of Ret isoforms. In cultured mammalian cells (HEK293 and SH-SY5Y cells), RET51 is preferentially localized to the plasma membrane, more rapidly internalized upon ligand binding and is differentially ubiquitylated compared to RET9 (Crupi et al., 2015; Hyndman et al., 2017). Different rates of Ret51 degradation are also seen in distinct neuronal populations. In mouse sympathetic neuron culture, RET51 is rapidly degraded in axon terminals following GDNF treatment (Tsui and Pierchala, 2010). However, in DRG sensory neuron cultures, RET51 was substantially more resistant to degradation and activated RET51 persisted much longer than in sympathetic neurons, which is correlated with detection of pRET51 in neuron cell bodies. Thus, it is possible that the plasma membrane localization and rapid internalization properties of Ret51 combined with the enhanced longevity of pRet51 in sensory neurons allow for its unique role in pioneer sensory neurons and, potentially, the ability to undergo long-range retrograde axonal transport.

In summary, our in vivo data demonstrate a novel role for Ret51 in pioneer axon extension and suggest the requirement of long-range, retrograde Ret signaling in this process. Additionally, we establish a system to examine the role of neurotrophic factor signaling in pioneer axon growth without a confounding role in neuronal survival. We identify that Jip3 mediates Ret51 retrograde transport, providing a useful tool to dissect local versus long-range signaling contributions of Ret receptor. Combined with our observation that Ret is specifically and highly expressed in pLLG pioneer sensory neurons, we submit that Ret51 functions as a molecular and transcriptional marker of these pioneers. This finding opens up several new technical avenues to examine the mechanisms underlying pioneer neuron development and pathfinding. In particular, this finding can be leveraged with transcriptomic techniques, such as single-cell RNA-seq, to study the transcriptional mechanisms that specify pioneer neurons and promote their enhanced axon growth and pathfinding capabilities. Finally, our results offer a holistic mechanism of long-range retrograde axonal transport of a neurotrophic factor receptor (Ret) inducing a specific transcriptional change (myo10l1) that ultimately impacts local protein levels in the growth cone (Myo10l1) required for proper axon behavior during development (filopodia formation and axon extension, Figure 7I). By elucidating the specific roles of concrete components involved in this process, our mechanistic account conceptually advance the paradigm of long-range neurotrophic factor signaling in axon growth. Better understanding of the fundamental processes that regulate the unique axonal growth and pathfinding capabilities of pioneer neurons can provide potential therapeutic points of intervention to promote axonal regeneration in response to injury or disease. Future studies will focus on identifying additional potential transcriptional targets downstream of Ret retrograde transport and how these regulate specific changes in pioneer growth cone morphology and behavior.

### Materials and methods

#### Key resources table

<table>
<thead>
<tr>
<th>Reagent type (species) or resource</th>
<th>Designation</th>
<th>Source or reference</th>
<th>Identifiers</th>
<th>Additional information</th>
</tr>
</thead>
<tbody>
<tr>
<td>Strain (Danio rerio)</td>
<td>TgBAC(neurod:EGFP)int; neurod:EGFP</td>
<td>DOI: 10.1523/JNEUROSCI.5230–07.2008</td>
<td>ZRC Catalog ID: ZL3218</td>
<td></td>
</tr>
<tr>
<td>Strain (Danio rerio)</td>
<td>rethu2866</td>
<td>DOI: 10.1242/dev.061002</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Strain (Danio rerio)</td>
<td>jip3nl7</td>
<td>DOI: 10.1371/journal.pgen.1003303</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Antibody</td>
<td>SCG10(STMN2) antibody (rabbit polyclonal)</td>
<td>ProteinTech Cat. No.: 10586-AP 1:100</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Antibody</td>
<td>GFP (rabbit polyclonal)</td>
<td>Invitrogen Cat. No.: A11122 1:1000</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Antibody</td>
<td>GFP (chick polyclonal)</td>
<td>Aves Labs Cat. No.: GFP-1020 1:1500</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

Continued on next page
<table>
<thead>
<tr>
<th>Reagent type (species) or resource</th>
<th>Designation</th>
<th>Source or reference</th>
<th>Identifiers</th>
<th>Additional information</th>
</tr>
</thead>
<tbody>
<tr>
<td>Antibody</td>
<td>Phospho-Ret (pTyr905) (rabbit polyclonal)</td>
<td>Invitrogen</td>
<td>Cat. No.: PA517761</td>
<td>1:50</td>
</tr>
<tr>
<td>Antibody</td>
<td>Ret (rabbit polyclonal)</td>
<td>Santa Cruz Biotechnology</td>
<td>Cat. No.: sc-167</td>
<td>1:100</td>
</tr>
<tr>
<td>Antibody</td>
<td>Neurofilament (mouse monoclonal)</td>
<td>Developmental Studies Hybridoma Bank</td>
<td>Designation: 3A10</td>
<td>1:500</td>
</tr>
<tr>
<td>Antibody</td>
<td>Myosin-X (rabbit polyclonal)</td>
<td>ProteinTech</td>
<td>Cat. No.: 24565-1-AP</td>
<td>1:500</td>
</tr>
<tr>
<td>Antibody</td>
<td>Phospho-JNK (Thr183/Tyr185) (rabbit polyclonal)</td>
<td>Cell Signaling</td>
<td>Cat. No.: 9251S</td>
<td>1:100</td>
</tr>
<tr>
<td>Antibody</td>
<td>Goat anti-chicken Alexa Fluor 488 secondary antibody</td>
<td>Invitrogen</td>
<td>Cat. No.: A-11039</td>
<td>1:10000</td>
</tr>
<tr>
<td>Antibody</td>
<td>Goat anti-rabbit Alexa Fluor 568 secondary antibody</td>
<td>Invitrogen</td>
<td>Cat. No.: A-11008</td>
<td>1:10000</td>
</tr>
<tr>
<td>Antibody</td>
<td>Goat anti-mouse Alexa Fluor 568 secondary antibody</td>
<td>Invitrogen</td>
<td>Cat. No.: A-11004</td>
<td>1:10000</td>
</tr>
<tr>
<td>Antibody</td>
<td>Cy5 Goat anti-mouse IgG secondary antibody</td>
<td>Jackson ImmunoResearch Laboratories</td>
<td>Cat. No.: 115-175-003</td>
<td>1:10000</td>
</tr>
<tr>
<td>Antibody</td>
<td>Goat anti-rabbit Alexa Fluor 647 secondary antibody</td>
<td>Jackson ImmunoResearch Laboratories</td>
<td>Cat. No.: A-27040</td>
<td>1:10000</td>
</tr>
<tr>
<td>Recombinant DNA reagent</td>
<td>pCDNA3.1-RET9-mCherry</td>
<td>DOI: 10.1111/tra.12315</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Recombinant DNA reagent</td>
<td>pCDNA3.1-RET51-mCherry</td>
<td>DOI: 10.1111/tra.12315</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Recombinant DNA reagent</td>
<td>neurod5kb:mCherry</td>
<td>This paper</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Recombinant DNA reagent</td>
<td>neurod5kb:RET51-mCherry</td>
<td>This paper</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Recombinant DNA reagent</td>
<td>TgBAC(ret51-eGFP)</td>
<td>This paper</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

Continued on next page
Zebrafish husbandry

Adult zebrafish were maintained at 28.5˚C as previously described and embryos were derived from natural matings or in vitro fertilization (Westerfield, 2000), raised in embryo media, and developmentally staged (Kimmel et al., 1995). Strains utilized were AB, TgBAC(neurod:EGFP)nl1 (Obholzer et al., 2008), rethu2846 (Knight et al., 2011), and jipnl7 (Drerup and Nechiporuk, 2013).

Plasmids and tagged constructs

Neurod5kb:jip3-mCherry was previously described (Drerup and Nechiporuk, 2013). Neurod5kb:mCherry plasmid was generated by recombining the neurod5kb promoter (Mo and Nicolson, 2011) p5e entry vector into the pDEST394 vector in the Gateway system by previously described methods (Kwan et al., 2007). Human RET9-mCherry and RET51-mCherry constructs in the EGFP-N1 vector were obtained from the Mulligan lab (Crupi et al., 2015) and digested with HindIII, NotI, and SphI to release the whole tagged construct and digest the EGFP-N1 vector backbone. Released RET fusion construct cassettes were ligated into HindIII/NotI-digested Gateway pME-MCS vector and subsequently recombined with the neurod5kb or cmv/sp6 promoter p5e entry vectors to yield neurod5kb:RET9/51-mCherry or cmv/sp6:RET9/51-mCherry in the pDEST394 destination vector (Kwan et al., 2007). Jip3 deletion constructs Δp150 and ΔJNK were derived from pME-Jip3 (Drerup and Nechiporuk, 2013) using the Quickchange II kit (Agilent) (primers in Table 1). mRNA was synthesized from Jip3 constructs or cmv/sp6:RET9/51-mCherry using SP6 mMessage Machine (Life Technologies) and microinjected at 500 pg/embryo for Jip3 constructs and 50 pg/embryo for RET9/51-mCherry constructs.

CRISPR-Cas9-mediated knockout

Three sgRNAs for CRISPR-Cas9 knockdown of myo10l1 were designed and injections were performed as previously described (Shah et al., 2015) (Table 1), with the modification that, instead of synthesizing and injecting individual sgRNAs, a cocktail of all three sgRNAs was amplified and synthesized simultaneously in one reaction (Wu et al., 2018), then injected in neurod:EGFP embryos. At 4 dpf, embryos with obvious necrosis or body morphology defects were removed and body segment location of pLLG axon terminals was scored for injected and uninjected embryos, and genomic DNA was isolated for individual embryos. CRISPR efficiency was evaluated by amplifying regions surrounding CRISPR cut sites (primers in Table 1) and incubating amplicons with restriction enzymes with cut sites adjacent to CRISPR targets (gRNA A: PstI; gRNA B: BssI; gRNA C: BstNI).

Generation of the Ret51- and Myo10l1-eGFP BAC fusions

We modified Ret- and Myo10l1-containing bacterial artificial chromosome (BAC) clones by Escherichia coli-based homologous recombination (Suster et al., 2011). BAC clone DKEY-192P21 contains 49.89 kb of sequence upstream and 69.78 kb of sequence downstream of ret; BAC clone DKEY-10E5 contains 20.98 kb sequence upstream and 48.7 kb of sequence downstream of myo10l1.
After recombination, the modified ret BAC clone contained an eGFP gene positioned in frame at the 3’ end of exon 20 to generate Ret51 fusion, following SV40 polyadenylation signal and kanamycin resistance gene. The modified Myo10l1 BAC clone contained an eGFP gene positioned in frame at the 3’ end of exon 40 to generate My10l1 fusion. The kanamycin resistance gene, flanked by FLP sites, was removed by FLP-mediated recombination (Suster et al., 2011). The accuracy of EGFP integration and FLP recombination were evaluated by PCR, sequencing, and by transient expression assays.

**In situ hybridization and wholemount immunostaining**

Templates for digoxygenin-labeled antisense RNA probes were generated for the ret9 specific region and ret9+51 by amplifying the alternatively spliced 3’ exons and 3’ UTR from cDNA using 3’ Rapid Amplification of cDNA Ends (3’-RACE) (Scotto-Lavino et al., 2006) with listed primers (Table 1). ‘3’RACE-d(T)’ primer was used for the initial reverse transcription reaction to generate cDNA specifically. ‘3’Anchor R’ primer specific to the end of 3’RACE-d(T) oligo was first used with a Ret-3’RACE-F primer to amplify both isoforms. This PCR product was subjected to a nested PCR with either ‘Ret9-3’unique RACE F’ or ‘Ret9+51 RACE F’ primers to amplify either a unique regions for ret9 or a region containing both ret9+51 probes, respectively, and cloned into pCR4-TOPO (Thermofisher). Fluorescence in situ hybridization was performed as previously described (Jülich et al., 2005) by digesting PCR4-TOPO clones with SpeI or NotI and digoxygenin-labeled...
antisense RNA probe synthesized with T7 or T3 polymerase (Life Technologies). *myo10l1* digoxigenin-labeled antisense RNA probe was generated using an EST containing 4.5 kb of *myo10l1* (Dharmacon, #EDR1052-208859531), digesting with Smal and synthesized with T7 polymerase (Life Technologies).

Whole mount immunohistochemistry was performed following established protocols (Ungos et al., 2003) with the following exception: embryos stained with anti-Ret were fixed in Shandon Glyo-Fixx (Thermo Scientific) for one hour at room temperature. The following antibodies were used: anti-GFP (1:1000; Invitrogen #A11122), anti-SCG10 (1:100; Proteintech, #10586-AP) anti-p905Ret (1:50; Thermo Pierce, #PA517761), anti-Ret (1:100; Santa Cruz, #sc-167), anti-neurofilament (1:500; DSHB, 3A10), anti-Myo10 (1:500; Proteintech #24565–1-AP), anti-pJNK (1:100; Cell Signaling #9251S). tRet antibody was validated by verifying lack of immunofluorescence signal in pLLG in ret
hu2864 mutant embryos compared to wild-type siblings (Figure 1—figure supplement 1, panels B, C). pRet antibody was validated by injecting a plasmid construct encoding RET51-mCherry (*neurod5kb:RET51-mCherry*), identifying embryos with 2–3 pLLG neurons expressing the construct, fixing at 30 hpf, and immunostaining. Co-localization of Ret51-mCherry and pRet was found in extending growth cones, but pRet was largely absent in the ganglion or cell bodies expressing Ret51-mCherry (Figure 1—figure supplement 1, panels D,E), consistent with expectation that only a subpopulation of Ret51 would be phosphorylated and that subpopulation would be enriched in extending growth cones near the source of GDNF ligand. For pioneer nerve sever experiments (Figure 3C–E), a-rabbit AlexaFluor 647 secondary (Thermo Fisher) was used to distinguish a-Ret antibody signal from rhodamine. All fluorescently labeled embryos were imaged using a 40X/NA = 1.3 oil objective on a FV1000 laser scanning confocal system (Olympus). Brightness and contrast were adjusted in Adobe Photoshop and figures were compiled in Adobe Illustrator.

**Quantification of immunofluorescence**

For analysis of pRet and tRet intensity in axon terminals and after nerve injury, individuals were immunolabeled as described above. For consistency of labeling, compared larvae were processed in the same batch. Confocal Z-stacks (0.5 μm between planes) were taken of the area of interest using a 40X/NA = 1.3 oil objective with identical settings. Images were analyzed using ImageJ (Abramoff et al., 2004). For fluorescence intensity measurements of pRet or tRet in wild type and mutant growth cones, summed projections of the regions of interest were generated only through regions that contained the *neurod:EGFP* signal and converted to eight bit in ImageJ as previously described (Drerup and Nechiporuk, 2013). Briefly, in the pLL nerve injury analysis, a 30 μm, *neurod:EGFP*-positive region encompassing the proximal or distal edge of the severed axon was selected and summed projections through only this segment were compiled for analysis. Prior to statistical comparison, the mean background fluorescent intensity, measured in a region adjacent to the injury site, was subtracted from the values generated.

For Myo10 immunostaining and *myo10l1* fluorescent in situ hybridization, wholemount embryos were imaged as described above using 488 and 568 nm excitation channels. Myo10 or *myo10l1* signal intensity was measured in pLL ganglia and collective axon terminals using Imaris (Bitplane) as follows. We used EGFP signal to generate surfaces of either the ganglia or the distal 50 μm of pioneer neuron terminals based on full stacks. We measured 568 nm channel mean fluorescence intensity of axon terminals or ganglia and subtracted the average mean fluorescence intensity of three regions proximal to the ganglion/axon terminals to account for background staining.

**Growth cone imaging and analysis**

For imaging, embryos were mounted in 1.5% low melting point agarose on a glass coverslip, submerged in embryo media containing 0.02% tricaine (MS-222; Sigma) and imaged at 24 or 30 hpf using a 60X/NA = 1.2 water objective on an upright Fluoview1000 confocal microscope (Olympus). The distal 75–125 μm of pLLG axon terminals, indicated by expression of *neurod:EGFP* transgene, were imaged with stacks of sufficient depth to capture all portions of the terminal axons. For individual growth cone imaging, embryos were injected with *neurod5kb:mCherry* plasmid, embryos expressing mCherry in 1–2 pLLG neurons with axons extending at or near the axon terminals (marked by *neurod:EGFP* expression) were selected and imaged as described using 488 and 568 nm excitation channels. Growth cone and collective axon terminal volume was quantified using Imaris.
software (Bitplane) as described above. EGFP signal was used to generate axonal surfaces of individual growth cones or the distal 75 μm of collective axon terminals, and volume was measured in Imaris. Filopodia were visualized by creating Z-projections of image stacks of individual growth cones in ImageJ and filopodia were traced using the segmented line tool, measured for length, and filopodia ≥1 μm number and length were scored.

**Retrograde axon labeling experiment**

40 hpf *neurod:EGFP* embryos were anesthetized and mounted in 1.2% low melt agarose. Tails were severed with rhodamine dextran-soaked scissors just rostral to pLLP growth cones. Embryos were freed from agarose, allowed to develop for 3 hr, then re-mounted and imaged using the FV1000 confocal microscope. Embryos were subsequently fixed and stained to detect Ret immunoreactivity.

**Axon transport analysis**

Zygotes were injected with plasmid DNA encoding fluoresently tagged cargos of interest with expression driven by the 5kbneurod promoter (Mo and Nicolson, 2011) and imaged as described above in ‘growth cone imaging and analysis.’ At 30 hpf, embryos were sorted under epifluorescence to identify individuals with tagged cargo expression in a few cells of the pLL ganglion. For each embryo, a region of interest (30–150 μm) was selected in the pLL nerve in which a long stretch of axon was observable in a single plane. Scans were taken at three frames per second for 500–1000 frames. Embryos were subsequently released from agarose and processed for genotyping. For co-transport, embryos expressing both constructs in a single cell were selected and imaged as described above using sequential imaging of the 488 and 568 nm excitation channels. 500 frames were collected at three frames per second.

Transport parameters were analyzed using kymograph analysis in the MetaMorph software package (Molecular Devices, Inc) as previously described (Drerup and Nechiporuk, 2016). Kymographs containing 10 or more traces were analyzed and each were averaged within individual embryos for statistical analysis. The number of particles moving in each direction was estimated based on traces on the kymographs and then normalized to length of axonal segment and total imaging time.

**Axotomy and image acquisition**

Five-day old zebrafish larvae (*neurod:EGFP* carriers) were anesthetized in 0.02% tricaine and embedded in 3% methylcellulose on a slide. Pulled thick-walled glass capillaries were used to sever the nerve between NMs 2 and 3. Slides were immersed in Ringer’s solution (116 mM NaCl, 2.9 mM KCl, 1.8 mM CaCl$_2$, 5 mM HEPES pH = 7.2, 1% Pen/Strep) and incubated at 28.5˚C for 3 hr. Larvae were then collected and immunolabeled for pRet or tRet and EGFP.

**Sorafenib drug treatment**

Sorafenib was dissolved in DMSO to make a 10 mM stock. Embryos were transferred at 24 hpf into embryo media containing 0.5% DMSO and 5 μM sorafenib. Embryos were placed into freshly prepared sorafenib/DMSO media at 48 hpf and then scored for axon truncation at 72 hpf.

**Statistical analysis**

Data were analyzed with the SPSS software package (IBM). Data suitable for parametric analysis were analyzed using ANOVA with post-hoc Tukey test. Data not suitable for parametric analysis were analyzed using Mann-Whitney U test (Wilcoxon ranked sum test). Axon truncation length measured within groups over time (Figure 1—figure supplement 2, panel A) was analyzed by repeated measures ANOVA with post hoc Bonferroni test for pairwise comparison. For experiments involving two independent variables (rescue of ret mutant phenotype with RET isoform mRNA injection or injection of myo10l1 gRNA/Cas9 in ret mutants and siblings) two-way ANOVA was performed to test if main effects and interactions were statistically significant. If interaction was statistically significant, ANOVA was repeated with simple effects, and significance of main effects was re-evaluated and post hoc tests were performed.
Ethics statement
All animal works were approved by and conducted according to guidelines of the Oregon Health and Science University IACUC.

Acknowledgements
We thank Dr. Lois Mulligan for her gift of RET9-mCherry and RET51-mCherry constructs and helpful discussion. Funding was provided to AVN from NICHD (1R01HD072844; http://www.nichd.nih.gov) and NINDS (1R01NS111419; http://www.ninds.nih.gov).

Additional information

<table>
<thead>
<tr>
<th>Funding</th>
<th>Grant reference number</th>
<th>Author</th>
</tr>
</thead>
<tbody>
<tr>
<td>Eunice Kennedy Shriver National Institute of Child Health and Human Development</td>
<td>1R01HD072844</td>
<td>Alex V Nechiporuk</td>
</tr>
<tr>
<td>National Institute of Neurological Disorders and Stroke</td>
<td>1R01NS111419</td>
<td>Alex V Nechiporuk</td>
</tr>
</tbody>
</table>

The funders had no role in study design, data collection and interpretation, or the decision to submit the work for publication.

Author contributions
Adam Tuttle, Conceptualization, Formal analysis, Validation, Investigation, Visualization, Methodology, Writing—original draft, Project administration, Writing—review and editing; Catherine M Drerup, Validation, Investigation, Methodology, Writing—review and editing; Molly Marra, Investigation, Methodology, Writing—review and editing; Hillary McGraw, Investigation, Writing—review and editing; Alex V Nechiporuk, Conceptualization, Resources, Supervision, Funding acquisition, Validation, Investigation, Visualization, Methodology, Writing—original draft, Project administration, Writing—review and editing

Author ORCID
Adam Tuttle https://orcid.org/0000-0002-4300-3848
Catherine M Drerup https://orcid.org/0000-0002-0219-3075
Alex V Nechiporuk https://orcid.org/0000-0002-8295-8188

Ethics
Animal experimentation: This study was performed in strict accordance with the recommendations in the Guide for the Care and Use of Laboratory Animals of the National Institutes of Health. All of the animals were handled according to approved institutional animal care and use committee (IACUC) protocols of Oregon Health and Science University. The protocol was approved by the IACUC committee of the OHSU Research Integrity Office (Protocol ID: IP00000495). All live imaging and larval and adult procedures were performed under MS-222 anesthesia, and every effort was made to minimize suffering.

Decision letter and Author response
Decision letter https://doi.org/10.7554/eLife.46092.028
Author response https://doi.org/10.7554/eLife.46092.029

Additional files
Supplementary files
• Transparent reporting form
Data availability
Experimental data underlying Figures 1–7 have been uploaded with the manuscript for publishing as source data.

References
Behili AB, Robertson BW, Cheney RE. 2006. Myosin-X is a molecular motor that functions in filopodia formation. PNAS 103:12411–12416. DOI: https://doi.org/10.1073/pnas.0602443103, PMID: 16894163
Drerup CM, Nechiporuk AV. 2013. JNK-interacting protein 3 mediates the retrograde transport of activated c-Jun N-terminal kinase and lysosomes. PLOS Genetics 9:e1003303. DOI: https://doi.org/10.1371/journal.pgen.1003303, PMID: 23468645
Gasanov EV, Rafieva LM, Korzh VP. 2015. BDNF-TrkB Axis regulates migration of the lateral line primordium and modulates the maintenance of mechanoreceptor progenitors. PLOS ONE 10:e0119711. DOI: https://doi.org/10.1371/journal.pone.0119711, PMID: 25751404


